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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/976,927	10/11/2001	G. Ramanath	020752-000111US	2055
20350	7590	11/24/2003	EXAMINER	
TOWNSEND AND TOWNSEND AND CREW, LLP TWO EMBARCADERO CENTER EIGHTH FLOOR SAN FRANCISCO, CA 94111-3834			KIELIN, ERIK J	
		ART UNIT	PAPER NUMBER	
		2813		

DATE MAILED: 11/24/2003

Please find below and/or attached an Office communication concerning this application or proceeding.

<b>Office Action Summary</b>	Application No.	Applicant(s)
	09/976,927	RAMANATH ET AL.
	Examiner	Art Unit
	Erik Kielin	2813

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --  
**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

1) Responsive to communication(s) filed on 11 September 2003.

2a) This action is **FINAL**.                    2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

4) Claim(s) 29-36 is/are pending in the application.

4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.

5) Claim(s) \_\_\_\_\_ is/are allowed.

6) Claim(s) 29-36 is/are rejected.

7) Claim(s) \_\_\_\_\_ is/are objected to.

8) Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on \_\_\_\_\_ is/are: a) accepted or b) objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. §§ 119 and 120**

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) All b) Some \* c) None of:

1. Certified copies of the priority documents have been received.

2. Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.

3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

13) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application) since a specific reference was included in the first sentence of the specification or in an Application Data Sheet. 37 CFR 1.78.

a) The translation of the foreign language provisional application has been received.

14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121 since a specific reference was included in the first sentence of the specification or in an Application Data Sheet. 37 CFR 1.78.

**Attachment(s)**

1)  Notice of References Cited (PTO-892)                    4)  Interview Summary (PTO-413) Paper No(s). \_\_\_\_\_ .

2)  Notice of Draftsperson's Patent Drawing Review (PTO-948)                    5)  Notice of Informal Patent Application (PTO-152)

3)  Information Disclosure Statement(s) (PTO-1449) Paper No(s) \_\_\_\_\_ .                    6)  Other: \_\_\_\_\_ .

## **DETAILED ACTION**

This action responds to the Amendment file 11 September 2003 (Paper no. 16).

### ***Information Disclosure Statement***

1. Examiner acknowledges Applicant's remarks in response to the objection to the IDS that an IDS had been filed about one month prior to the mailing of the first Office action on the merits. This IDS was not, however, matched with the application until after mailing of the action and Examiner was clearly unaware that such IDS existed. Examiner apologizes on behalf of the Office for any inconvenience.

Nonetheless, the information disclosure statement filed 18 July 2002 fails to comply with the provisions of 37 CFR 1.97, 1.98 and MPEP § 609 because some of the references have not been provided with dates in accordance with 37 CFR 1.98(b)(5). Also the MPEP 609 states,

**"Each publication must be identified by publisher, author (if any), title, relevant pages of the publication, and date and place of publication. The date of publication supplied must include at least the month and year of publication, except that the year of publication (without the month) will be accepted if the applicant points out in the information disclosure statement that the year of publication is sufficiently earlier than the effective U.S. filing date and any foreign priority date so that the particular month of publication is not in issue."** (Emphasis added.)

The IDS has been placed in the application file, but only the references initialed by Examiner have been considered. Applicant is advised that the date of any re-submission of any item of information contained in this information disclosure statement or the submission of any missing element(s) will be the date of submission for purposes of determining compliance with the

requirements based on the time of filing the statement, including all certification requirements for statements under 37 CFR 1.97(e). See MPEP § 609 ¶ C(1).

This is repeated from the action filed 26 February 2003.

***Claim Rejections - 35 USC § 103***

2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

3. Claims 29-33 and 36 are rejected under 35 U.S.C. 103(a) as being unpatentable over US 5,079,600 (**Schnur et al.**) in view of ASM Handbook Vol. 5, Surface Engineering, ASM International: Materials Park, Ohio, 1994, pp. 315-318, and the basic textbook by **Porterfield, Inorganic Chemistry, A Unified Approach**, Addison-Wesley: Reading, Massachusetts, 1984, pp. 487-488.

Regarding claim 29, **Schnur** discloses a semiconductor device comprising,

(a) a substrate (Fig. 1A);  
(b) a diffusion barrier (Fig. 1A, called “thin film”), wherein the diffusion barrier comprises a self-assembled monolayer, SAM, (col. 10, lines 42-47) including a plurality of molecules, each molecule having an aromatic group at the terminus of the molecule. (For example, EXAMPLE 28 at col. 21, discloses a SAM formed from trichloro-(4-pyridyl)-ethyl-silane which forms the equivalent SAM barrier layer as shown in structure entitled “1” on p. 7 of the instant specification. The pyridyl is an aromatic group.

(c) a metal layer (called “catalyst” and “metal layer”) on the diffusion barrier.

In pertinent part, **Schnur** states in col. 20,

**“EXAMPLE 24**

“Fabrication of MOS capacitor test structures.

“An n-type silicon wafer with a 100 nm thick thermal oxide layer was cleaned and treated with UTF3 as in example 14. The film was patterned using a mask with standard capacitor test structures and irradiated for 28 minutes with an Hg/Ar lamp. The wafer was metallized with the standard copper plating procedures, used in Example 5, yielding metal squares 800 microns on a side (area=6x10<sup>-3</sup> cm<sup>2</sup>). The metal/thermal oxide/n-type silicon (MOS) capacitors were then characterized by probing the metal pads and the back of the wafer with a Micromanipulator automatic C-V measuring system. The capacitance was found to be 26 pF/cm<sup>2</sup> with minimal (10 mV) hysteresis and remained stable at room temperature for at over 3 weeks, indicating that device degradation due to masked metal contamination (**diffusion of copper into the thermal oxide**) was not a problem. (Emphasis added.)

Accordingly, it is seen to be inherent that the SAM of **Schnur** is a diffusion barrier because **Schnur** states that “diffusion of copper into the thermal oxide” does not occur.

(See also section entitled, “Summary of the Invention”) the sections entitled “EXAMPLE 1” col. 11, lines 24-58 wherein the barrier layer is formed from “octenyldimethylchlorosilane” covalently bonded to the substrate. See also “EXAMPLE 3” and “EXAMPLE 5.”)

**Schnur** does not teach that the catalyst is copper, thereby having the limitation that “for each molecule of the plurality of molecules, the copper in the metal layer is in direct contact with the aromatic group of the molecule.”

**ASM Handbook** teaches that copper may be used as a catalyst for electroless plating of copper (pp. 315-318 --especially p. 318 sections entitled, “Catalyzation” and “Copper catalyst”). The basic textbook of **Porterfield** ensures that copper forms metal complexes with pyridine

groups, such as the pyridine group used in **Schnur** as the polar end-group of each molecule in the SAM barrier layer.

It would have been obvious for one of ordinary skill in the art, at the time of the invention to use copper as the metal catalyst in **Schnur** as taught in the **ASM Handbook** because **Schnur** is not limited to Pd/Sn catalysts, as at least claim 1 of **Schnur** makes clear, and because copper is a known catalyst for electroless copper plating, as used in **Schnur**, as taught by the **ASM Handbook**. In this regard, it has been held that the selection of a known material based upon its suitability for an intended purpose is obvious. Moreover, the **ASM Handbook** teaches that copper-based catalyst are less expensive than palladium-based catalyst, thereby providing additional motivation to use copper for cost savings.

**Schnur** only requires that the catalyst bond to polar end group of the molecule which is the pyridyl end group of “Example 28” in **Schnur**. **Porterfield** ensures that such bonding occurs, such the one of ordinary skill has a reasonable expectation of success for using copper instead of Pd/Sn, as the catalyst in **Schnur**. Accordingly, using copper as the catalyst gives the limitation, that “for each molecule of the plurality of molecules, the copper in the metal layer is in direct contact with the aromatic group of the molecule.”

Regarding claim 30, it is held absent evidence to the contrary, that the diffusion barrier of **Schnur** is capable of preventing the diffusion of metal atoms from the metal layer into the substrate when the semiconductor device is exposed to thermal annealing at 200 °C or an electric field of 2 MV/cm at 200 °C in flowing N<sub>2</sub>. Basis for this reasoning is that Applicant is using the exact same SAM, as is **Schnur** to form the barrier layer. **Schnur** also points out in Example 24 that copper diffusion does not occur even under the stress of an electric field. See *In re Swinhart*,

169 USPQ 226,229 (CCPA 1971) (where the Patent Office has reason to believe that a functional limitation asserted to be critical for establishing novelty in the claimed subject matter may, in fact, be an inherent characteristic of the prior art, it possesses the authority to require the applicant to prove that subject matter shown to be in the prior art does not possess the characteristics relied on) and *In re Fitzgerald*, 205 USPQ 594 (CCPA 1980) (the burden of proof can be shifted to the applicant to show that subject matter of the prior art does not possess the characteristic relied on whether the rejection is based on inherency under 35 USC 102 or obviousness under 35 USC 103). Note that as long as there is evidence of record establishing inherency, failure of those skilled in the art to contemporaneously recognize an inherent property, function or ingredient of a prior art reference does not preclude a finding of anticipation. *Atlas Powder Co. v. IRECO, Inc.*, 190 F.3d 1342, 1349, 51 USPQ2d 1943, 1948 (Fed. Cir. 1999) (Two prior art references disclosed blasting compositions containing water-in-oil emulsions with identical ingredients to those claimed, in overlapping ranges with the claimed composition. The only element of the claims arguably not present in the prior art compositions was “sufficient aeration . . . entrapped to enhance sensitivity to a substantial degree.” The Federal Circuit found that the emulsions described in both references would inevitably and inherently have “sufficient aeration” to sensitize the compound in the claimed ranges based on the evidence of record (including test data and expert testimony). This finding of inherency was not defeated by the fact that one of the references taught away from air entrapment or purposeful aeration.). See also *In re King*, 801 F.2d 1324, 1327, 231 USPQ 136, 139 (Fed. Cir. 1986); *Titanium Metals Corp. v. Banner*, 778 F.2d 775, 782, 227 USPQ 773, 778 (Fed. Cir. 1985).

Regarding claim 31, as noted above, Example 24 in **Schnur** states that the substrate is a silicon wafer with silicon oxide formed thereon.

Regarding claim 32, the first end of the **Schnur** molecule is the pyridyl group, which is an aromatic group.

Regarding claim 33, the method by which the metal layer is deposited does not have patentable weight. Note that a “product by process” claim is directed to the product *per se*, no matter how actually made, *In re Hirao*, 190 USPQ 15 at 17 (footnote 3). See also *In re Brown*, 173 USPQ 685; *In re Luck*, 177 USPQ 523; *In re Fessmann*, 180 USPQ 324; *In re Avery*, 186 USPQ 161; *In re Wertheim*, 191 USPQ 90 (209 USPQ 554 does not deal with this issue); *In re Marosi* et al, 218 USPQ 289; and particularly *In re Thorpe*, 227 USPQ 964, all of which make it clear that it is the patentability of the final product *per se* which must be determined in a “product by process” claim, and not the patentability of the process, and that an old or obvious product produced by a new method is not patentable as a product, whether claimed in “product by process” claims or not. Note that applicant has the burden of proof in such cases, as the above case law make clear.

4. Claims 34 and 35 are rejected under 35 U.S.C. 103(a) as being unpatentable over US 5,079,600 (**Schnur** et al.) in view of **ASM Handbook** and **Porterfield**, as applied to claim 29 above, and considered with the article **Simon** et al. “Synthesis and characterization of a new surface derivitizing reagent to promote the adhesion of polypyrrole films to n-type silicon photoanodes: N-(3-(trimethoxysilyl) propyl)pyrrole” Journal of the American Chemical Society, 104, Dec. 1982, pp. 2031-2034 for a showing of inherency only.

Regarding claim 34, **Schnur** does not show how that SAM is bonded to the silicon oxide layer via three oxygen atoms, but this is inherent.

For verification that the trimethoxysilyl group inherently leads to the three Si-O bonds as instantly claimed, see the article **Simon** et al. "Synthesis and characterization of a new surface derivitizing reagent to promote the adhesion of polypyrrole films to n-type silicon photoanodes: N-(3-(trimethoxysilyl) propyl)pyrrole" Journal of the American Chemical Society, 104, Dec. 1982, pp. 2031-2034 --especially see the second page, the left-hand column, and Fig. 1 in the right-hand column.)

Regarding claim 35, **Schnur** discloses that the SAM is formed from the claimed molecule. **Schnur** uses the silane compound  $\beta$ -trichloro(4-pyridyl) ethyl silane to form the SAM in "Example 28." The "ethyl" syllable refers to the  $(CH_2)_2$ - group, the silyl to the Si, and the pyridyl to the six-atom cyclic aromatic group containing the nitrogen wherein  $R^3$  and  $R^4$  are hydrogen. The trichloro refers to  $Cl_3Si$ - group which, upon reaction, produces the Si bonded to the 3 oxygen atoms which are, in turn, bonded near the surface of the substrate. The chloro groups serve the same function as do the methoxy groups in bonding the compound to the oxide surface.

#### *Response to Arguments*

5. Applicant's arguments filed 11 September 2003 have been fully considered but they are not persuasive.

Applicant argues that the ASM Handbook teaches away from the use of copper catalysts. Examiner respectfully but emphatically disagrees for two reasons. While Examiner

acknowledges that the ASM Handbook indicates that there exists a “major disadvantage,” the ASM Handbook in the two sentences directly preceding that sentence quoted from the ASM Handbook by Applicant states,

*“Copper catalyst: Because palladium is more expensive than the less noble metals (such as copper and nickel) that are known to be autocatalytic, there have been numerous attempts to develop a viable catalyst based on one of these metals. A copper-tin colloidal catalyst was first reported in 1976 (Ref 59) and has proven commercially successful in a number of installations.”* (Italicized emphasis in original; bold emphasis added.)

Accordingly, not only has the “major disadvantage” apparently been successfully avoided -- owing to the proven commercial success of copper-tin catalyst-- one of ordinary skill gains the added benefit of a less expensive material than palladium which will save money, as is always highly desired in the highly competitive art of semiconductor device manufacturing.

For this reason, it is improper for Applicant to argue,

“There is **no motivation** to modify Schnur et al. with the teachings of ASM, because the cited references explicitly teach away from the modification proposed by the Examiner.” (Emphasis added.)

This statement is factually in error given that the ASM Handbook suggests the proposed modification to save money and that copper-based catalysts have been proven commercially successful. Rather Applicant has ignored the express teachings of the ASM Handbook and has taken a single sentence out of context as if that were all the ASM Handbook were teaching.

The article provided by Applicant is not germane to the instant case because the metal being deposited is a Fe-Ni alloy --not copper. Moreover, the film is not being deposited on the given seed layer. Absent evidence that sputter deposited copper films and electroplated copper films on the Schnur barrier layer are different, the evidence will not be considered persuasive.

Finally it is noted that the inventive feature of the instant invention is the use of the self-assembled monolayer as barrier layer. The manner in which the copper is depositing has not been indicated to be critical to the instant invention and is not considered novel. (See instant specification p. 2, section entitled, "Summary of the Invention.") Schnur discloses the exact same barrier layer as is presently being claimed.

***Conclusion***

6. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Erik Kielin whose telephone number is 703-306-5980. The examiner can normally be reached on 9:00 - 19:30 on Monday through Thursday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Carl Whitehead, Jr., can be reached at 703-308-4940. The fax phone numbers for the organization where this application or proceeding is assigned are 703-872-9318 for regular communications and 703-872-9319 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0956.



Erik Kielin  
Primary Examiner  
November 22, 2003